

## CLAIMS:

- SUB A1
- (1) A recombinant MVA containing and capable of expressing one or more DNA sequences encoding dengue virus antigens.
- (2) A recombinant MVA according to claim 1 containing and capable of expressing DNA sequences encoding antigens from all four dengue virus serotypes (type 1, 2, 3 and 4).
- SUB A2 CONT'D.
- (3) A recombinant MVA according to claims 1 to 2, wherein the dengue virus antigen is selected from preM, E and/or NS1 antigens.
- (4) A recombinant MVA according to claims 1 to 3, wherein the DNA sequences are inserted at the site of naturally occurring deletions within the MVA genome.
- (5) A recombinant MVA according to claims 1 to 4, wherein the DNA sequences encoding antigen is under transcriptional control of the vaccinia virus early/late promoter P7.5.
- (6) A vaccine containing at least one recombinant MVA according to claims 1 to 5 and a pharmaceutically acceptable carrier or diluent.
- (7) A vaccine according to claim 6 containing a recombinant MVA encoding a dengue virus type 1 antigen; a recombinant MVA encoding a dengue virus type 2 antigen; a recombinant MVA encoding a dengue virus type 3 antigen, and/or a recombinant MVA encoding a dengue virus type 4 antigen, and a pharmaceutically acceptable carrier or diluent.
- (8) A method for the treatment or prevention of dengue virus infection comprising administering to a living animal body, including a human, in need thereof a therapeutically effective amount of a recombinant MVA according to claims 1 to 5, or a vaccine according to claims 6 to 7.
- (9) A vaccine comprising as a first component a recombinant MVA carrying and capable of expressing T7 RNA polymerase and as further components one or more recombinant DNA vectors each carrying at least one
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~~dengue virus antigen under transcriptional control of a T7 RNA polymerase promoter.~~

(10) A method for the treatment or prevention of a dengue virus infection comprising inoculating a living animal body, including a human, in need thereof with the first and further components of a vaccine according to claim 9 either simultaneously or with a timelag but using the same inoculation site.

sub A2  
conclude

ADD A3

add B

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Ca Add claims  
32-38